WEST VIRGINIA SECRETARY OF STATE

MAC WARNER

ADMINISTRATIVE LAW DIVISION

eFILED

4/4/2022 9:59:28 AM

Office of West Virginia Secretary Of State

NOTICE OF AGENCY APPROVAL OF A PROPOSED RULE AND FILING WITH THE LEGISLATIVE RULE-MAKING REVIEW COMMITTEE

AGENCY:

Pharmacy

TITLE-SERIES:

15-19

RULE TYPE:

Legislative

Amendment to Existing Rule:

Repeal of existing rule:

RULE NAME:

15-19 West Virginia Board Of Pharmacy Inspections

PRIMARY CONTACT

NAME:

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CITE STATUTORY AUTHORITY:

30-5-7

EXPLANATION OF THE STATUTORY AUTHORITY FOR THE LEGISLATIVE RULE, INCLUDING A DETAILED SUMMARY OF THE EFFECT OF EACH PROVISION OF THE LEGISLATIVE RULE WITH CITATION TO THE SPECIFIC STATUTORY PROVISION WHICH EMPOWERS THE AGENCY TO ENACT SUCH RULE PROVISION:

This rule governs inspections. The Board was required to file this emergency rule pursuant to a legislative audit.

IS THIS FILING SOLELY FOR THE SUNSET PROVISION REQUIREMENTS IN W. VA. CODE §29A-3-19(e)? No

IF YES, DO YOU CERTIFY THAT THE ONLY CHANGES TO THE RULE ARE THE FILING DATE, EFFECTIVE DATE AND AN EXTENSION OF THE SUNSET DATE? No

DATE eFiled FOR NOTICE OF HEARING OR PUBLIC COMMENT PERIOD:

2/3/2022

DATE OF PUBLIC HEARING(S) OR PUBLIC COMMENT PERIOD ENDED:

3/7/2022

COMMENTS RECEIVED:

Yes

(IF YES, PLEASE UPLOAD IN THE COMMENTS RECEIVED FIELD COMMENTS RECEIVED AND RESPONSES TO COMMENTS)

PUBLIC HEARING:

No

TRANSCRIPTS)
RELEVANT FEDERAL STATUTES OR REGULATIONS: No
WHAT OTHER NOTICE, INCLUDING ADVERTISING, DID YOU GIVE OF THE HEARING?
This rule was posted on the West Virginia Secretary of State's website.
SUMMARY OF THE CONTENT OF THE LEGISLATIVE RULE, AND A DETAILED DESCRIPTION OF THE RULE'S PURPOSE AND ALL PROPOSED CHANGES TO THE RULE:
This rule governs inspections by the West Virginia Board of Pharmacy.
STATEMENT OF CIRCUMSTANCES WHICH REQUIRE THE RULE:
This rule was required pursuant to the findings of a legislative audit.
SUMMARIZE IN A CLEAR AND CONCISE MANNER THE OVERALL ECONOMIC IMPACT OF THE PROPOSED LEGISLATIVE RULE:
A. ECONOMIC IMPACT ON REVENUES OF STATE GOVERNMENT:
none
B. ECONOMIC IMPACT ON SPECIAL REVENUE ACCOUNTS:
none
C. ECONOMIC IMPACT OF THE LEGISLATIVE RULE ON THE STATE OR ITS RESIDENTS:
none

(IF YES, PLEASE UPLOAD IN THE PUBLIC HEARING FIELD PERSONS WHO APPEARED AT THE HEARING(S) AND

D. FISCAL NOTE DETAIL:

Effect of Proposal	Fiscal Year			
	2022 Increase/Decrease (use "-")	2023 Increase/Decrease (use "-")	Fiscal Year (Upon Full Implementation)	
1. Estimated Total Cost	0	0	0	
Personal Services	O	0	0	
Current Expenses	0	0	0	
Repairs and Alterations	0	0	0	
Assets	0	0	0	
Other	0	0	0	
2. Estimated Total Revenues	0	0	0	

E. EXPLANATION OF ABOVE ESTIMATES (INCLUDING LONG-RANGE EFFECT):

none

BY CHOOSING 'YES', I ATTEST THAT THE PREVIOUS STATEMENT IS TRUE AND CORRECT.

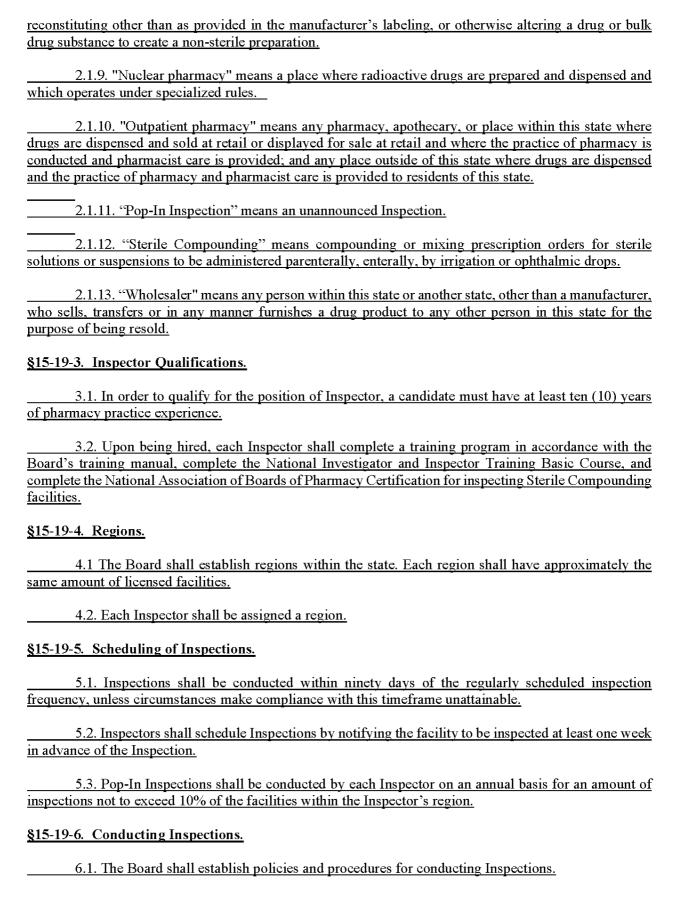
Yes

Ryan L Hatfield -- By my signature, I certify that I am the person authorized to file legislative rules, in accordance with West Virginia Code §29A-3-11 and §39A-3-2.

TITLE 15 LEGISLATIVE RULE WEST VIRGINIA BOARD OF PHARMACY

SERIES 19 INSPECTIONS

<u>§15-19-1. General.</u>
1.1. Scope Inspection process for the West Virginia Board of Pharmacy.
1.2. Authority W. Va. Code §§ 30-5-7.
1.3. Filing date
1.4. Effective date
1.5. Sunset Provision This rule will terminate and have no further force or effect on August 1, 2027.
§15-19-2. Definitions.
2.1. The following words and phrases as used in this Rule mean:
2.1.1. "Board" means the West Virginia Board of Pharmacy.
2.1.2. "Charitable Clinic Pharmacy" means a clinic or facility organized as a not-for-profit corporation that offers pharmaceutical care and dispenses prescriptions free of charge to appropriately screened and qualified patients. A charitable clinic pharmacy shall meet the minimum standards for a pharmacy as set forth in W. Va. Code §30-5-1, et seq., and by this rule, but may not be charged any applicable licensing fees. A charitable clinic pharmacy may have pharmacists-in-charge, as that term is defined in this section, who volunteers his or her services. A charitable clinic may also receive donated drugs. It is not the intent of this rule to affect any organizations which are merely operating a prescribing practitioner's or clinic's free sample drug room.
2.1.3. "Controlled Substance Permit" means the permit require to be obtained by the Board, unless otherwise exempt, for every person who manufactures, distributes, including reverse distributing, or dispenses any controlled substances or who proposes to engage in the manufacture, distribution or dispensing of any controlled substance within this state.
2.1.4. "Inspection" means the process by which the Board verifies certain information of Board licenses.
2.1.5. "Inspector" means a person employed by the Board to perform inspections.
2.1.6. "Institutional Pharmacy" means that physical portion of an institutional facility that is engaged in the compounding, dispensing, and distribution of drugs, devices, and other materials used in the diagnosis and treatment of injury, illness, and disease and which holds a pharmacy license from the Board. 2.1.7. "Non-compliance report" means a report created noting a deficiency discovered during an
inspection.
2.1.8 "Non-Sterile Compounding" means the process of combining admixing diluting pooling



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6.2. At a minimum, the methods established by the Board for conducting inspections shall include the following:
6.2.1. presentation of credentials;
6.2.2. copies of the inspection forms to be used:
6.2.3. details on the use of electronic inspection forms;
6.2.4. a listing of documents to be reviewed; and
6.2.5. listing of staff that may be interviewed.
6.3. Completed hard copy inspections shall be submitted to the Board office in a timely manner.
6.4. Completed hard copy inspections shall be recorded in the Board database.
6.5. Electronic inspection shall be automatically entered into the Board database.
6.6. Copies of the completed inspection form shall be provided to the facility inspected.
6.7. Non-compliance reports shall be completed and entered into the Board database.
6.8. Completed inspections shall be reviewed by the Chief Compliance Officer within ninety days. This review shall be documented in the Board database.
6.9. Non-compliance reports shall be logged in a manner which allows for the systematic monitoring of facility's actions toward correcting areas of significant deficiencies.
6.10. Facilities not meeting the expected corrections within the timeframe established by the inspector may be referred to the Complaint Committee of the Board.
6.11. All records, documents, communications, and images shall be maintained in the Board database for a period of five years.
§15-19-7. Inspection Forms.
7.1. The Board shall create inspection forms for each type of facility inspected. The inspection forms shall contain citations to relevant laws and regulations.
7.2. The inspection forms used by the Board shall be made available to the public on the Board's website.
§15-19-8. Inspection Frequency.
8.1. The following facility types shall be inspected annually:
8.1.1. outpatient pharmacies:
8.1.2. institutional pharmacies;
8.1.3. charitable clinic pharmacies;

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8.1.4. nuclear pharmacies:	
8.1.5. hazardous drug handling permit;	
8.1.6. sterile compounding facilities;	
8.1.7. non-sterile compounding facilities.	
8.2. Facilities that only hold a controlled substance permit and no other Board permit of be inspected biennially.	or license shal